CLAIMS

5

A method for rescuing damaged nerve cells in a patient, comprising:

 administering to a patient having damaged nerve cells an amount of a deprenyl
 compound such that rescuing of damaged nerve cells occurs in the patient;

with the proviso that the deprenyl compound is not selected from the group consisting of deprenyl, pargyline, AGN-1133, or AGN1135.

The method of claim 1, wherein the deprenyl compound is represented by the
 structure:

$$R_4-R_3-CH-N$$
 R_2
 R_5-R_6

in which

 R_1 is hydrogen, alkyl, alkenyl, alkynyl, aralkyl, alkylcarbonyl, arylcarbonyl, alkoxycarbonyl, or aryloxycarbonyl;

R2 is hydrogen or alkyl;

 R_3 is a single bond, alkylene, or -(CH₂)_n-X-(CH₂)_m;

in which X is O, S, or N-methyl; m is 1 or 2; and n is 0,1, or 2;

R4 is alkyl, alkenyl, alkynyl, heterocyclyl, aryl or aralkyl; and

R5 is alkylene, alkenylene, alkynylene and alkoxylene; and

R6 is C3-C6 cycloalkyl or

R₂ and R₄-R₃ are joined to form, together with the methine to which they are attached, a cyclic or polycyclic group;

and pharmaceutically acceptable salts thereof.

15

20

10

20

- 3. The method of claim 2, wherein R₁ is a group that can be removed in vivo.
- 4. The method of claim 2, wherein R₁ is hydrogen.
- 5 5. The method of claim 2, wherein R₁ is alkyl.
 - 6. The method of claim 5, wherein R1 is methyl.
 - 7. The method of claim 2, wherein R₂ is methyl.
 - 8. The method of claim 2, wherein R₃ is methylene.
 - 9. The method of claim 2, wherein R4 is aryl.
- 15 10. The method of claim 2, wherein R4 is phenyl.
 - 11. The method of claim 2, wherein R5 is methylene.
 - 12. The method of claim 2, wherein R6 is

—C≡CH

13. The method of claim 2, wherein the deprenyl compound has the structure

wherein R_1 is hydrogen, alkyl, alkenyl, alkynyl, aralkyl, alkylcarbonyl, arylcarbonyl, alkoxycarbonyl, or aryloxycarbonyl.

14. The method of claim 2. wherein the deprenyl compound is represented by the structure:

$$R_4-R_3-CH-N$$
 $R_3-CH_2-C \equiv CH$

in which

10

15

20

 R_1 is hydrogen, alkyl, alkenyl, alkynyl, aralkyl, alkylcarbonyl, arylcarbonyl, alkoxycarbonyl, or aryloxycarbonyl;

R2 is hydrogen or alkyl;

R3 is a bond or methylene; and

R4 is aryl or aralkyl; or

R₂ and R₄-R₃ are joined to form, together with the methine to which they are attached, a cyclic or polycyclic group;

and pharmaceutically acceptable salts thereof.

15. The method of claim 2, wherein the deprenyl compound is represented by the structure:

$$R_4-R_3-CH-N$$
 R_2
 $R_5-C=CH$

in which

R2 is hydrogen or alkyl;

R3 is a bond or methylene; and

R4 is aryl or aralkyl; or

 R_2 and R_4 - R_3 are joined to form, together with the methine to which they are attached, a cyclic or polycyclic group; and

R₅ is alkylene, alkenylene, alkynylene and alkoxylene; and pharmaceutically acceptable salts thereof.

16. The method of claim 2, wherein the deprenyl compound is represented by the structure:

$$\begin{array}{c} A_{n} \\ CH_{2}-CH-N \\ CH_{1} CH_{2}-C = CH \end{array}$$

in which

 R_1 is hydrogen, alkyl, alkenyl, alkynyl, aralkyl, alkylcarbonyl, arylcarbonyl, alkoxycarbonyl, or aryloxycarbonyl;

A is a substituent independently selected for each occurrence from the group consisting of halogen, hydroxyl, alkyl, alkoxyl, cyano, nitro, amino, carboxyl, -CF3, or azido;

n is 0 or an integer from 1 to 5; and pharmaceutically acceptable salts thereof.

- 17. The method of claim 1, wherein the deprenyl compound is (-)-desmethyldeprenyl.
- 18. A kit comprising a container of a deprenyl compound and instructions for administering a therapeutically effective amount of the deprenyl compound to a subject having damaged nerve cells such that rescuing of damaged nerve cells occurs in the subject.

10

15

20